IN THE CLAIMS

Please amend the claims as follows:

1.- (Currently Amended) A sulfonamide of general formula (Ia),

wherein

R¹ represents a –NR⁷R⁸ radical or a saturated or unsaturated, optionally at least monosubstituted, optionally at least one heteroatom as a ring member containing cycloaliphatic radical, which may be condensed with a saturated or unsaturated, optionally at least mono-substituted, optionally at least one heteroatom as a ring member containing mono- or bicyclic cycloaliphatic ring system,

R², R³, R⁴, R⁵ and R⁶, identical or different, each represent hydrogen, halogen, cyano, nitro, a saturated or unsaturated, linear or branched aliphatic radical, a linear or branched alkoxy radical, a linear or branched alkylthio radical, hydroxy, trifluoromethyl, a saturated or unsaturated cycloaliphatic radical, an alkylcarbonyl radical, a phenylcarbonyl or a –NR⁹R¹⁰ group,

R⁷ and R⁸, identical or different, each represent hydrogen or a saturated or unsaturated, optionally at least mono-substituted linear or branched aliphatic radical,

with the proviso that R⁸ and R⁹ are not hydrogen at the same time, and if one of them,

 R^8 or R^9 , is a saturated or unsaturated, linear or branched, optionally at least monosubstituted C_1 - C_4 aliphatic radical, the other one is a saturated or unsaturated, linear or branched, optionally at least mono-substituted aliphatic radical with at least five carbon atoms,

or

R⁷ and R⁸, together with the bridging nitrogen atom form a saturated or unsaturated, optionally at least mono-substituted, optionally at least one further heteroatom as a ring member containing heterocyclic ring which may be condensed with a saturated or unsaturated, optionally at least mono-substituted, optionally at least one heteroatom as a ring member containing mono- or bicyclic cycloaliphatic ring system,

R⁹ and R¹⁰, identical or different, each represent hydrogen or a saturated or unsaturated, linear or branched, optionally at least mono-substituted aliphatic radical,

or

R⁹ and R¹⁰, together with the bridging nitrogen atom form a saturated or unsaturated, optionally at least mono-substituted, optionally at least one further heteroatom as a ring member containing heterocyclic ring which may be condensed with a saturated or unsaturated, optionally at least mono-substituted, optionally at least one heteroatom as a ring member containing mono- or bicyclic cycloaliphatic ring system,

A and B, identical or different, each represent a saturated or unsaturated, linear or branched aliphatic radical, optionally at least mono-substituted

or

A and B, together with the carbon atom to which they are bonded, form a saturated or unsaturated, but not aromatic cycloalkyl ring, optionally at least mono-substituted

and

n is 0,

-a stereoisomer thereof, an enantiomer thereof, a diasteromer thereof, a racemate thereof, or a pharmaceutically acceptable salt thereof, or mixtures thereof.

- 2. (Previously Presented) The compound according to claim 1, wherein R¹ represents a -NR⁷R⁸ radical or a saturated or unsaturated, optionally at least monosubstituted, optionally at least one heteroatom as a ring member containing 5- or 6-membered cycloaliphatic radical, which may be condensed with a saturated or unsaturated, optionally at least mono-substituted, optionally at least one heteroatom as a ring member containing mono- or bicyclic cycloaliphatic ring system, whereby the rings of the ring system are 5- or 6-membered.
- 3.- (Previously Presented) The compound according to claim 1 wherein R², R³, R⁴, R⁵ and R⁶, identical or different, each represent hydrogen, F, Cl, Br, cyano, nitro, a linear or branched C₁₋₆ alkyl radical, a linear or branched C₂₋₆ alkenyl radical, a linear or branched C₁₋₆ alkoxy, a linear or branched C₁₋₆ alkylthio, hydroxy, trifluoromethyl, a saturated or unsaturated C₃₋₈ cycloaliphatic radical, a linear or branched C₁₋₆ alkylcarbonyl radical, phenylcarbonyl or an –NR⁹R¹⁰ group.
- 4.- (Previously Presented) The compound according to claim 1, wherein R^7 and R^8 , identical or different, each represent hydrogen, a linear or branched, optionally at least mono-substituted C_{1-10} alkyl radical, a linear or branched, optionally at least mono-substituted, C_{2-10} alkenyl radical, or a linear or branched, optionally at least mono-substituted, C_{2-10} alkynyl radical or

R⁷ and R⁸, together with the bridging nitrogen form a saturated or unsaturated, optionally at least mono-substituted, optionally at least one further heteroatom as a ring member containing 5- or 6-membered heterocyclic ring which may be condensed with a saturated or unsaturated, optionally at least mono-substituted, optionally at least one heteroatom as a ring member containing mono- or bicyclic cycloaliphatic ring system, whereby the rings of the ring system are 5- 6- or 7-membered.

5.- (Previously Presented) The compound according to claim 4, wherein R⁷ and R⁸, identical or different, each represent hydrogen or a linear or branched C₁₋₁₀ alkyl radical or

R⁷ and R⁸, together with the bridging nitrogen atom form a radical chosen from the group consisting of

wherein R²⁰, if present, is hydrogen, a linear or branched C₁-C₆ alkyl radical or a benzyl radical.

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- 6.- (Previously Presented) The compound according to claim 1, wherein R⁹ and R¹⁰, identical or different, each represent hydrogen, a linear or branched, optionally at least mono-substituted C₁-C₁₀ alkyl radical, a linear or branched, optionally at least mono-substituted C₂-C₁₀ alkenyl radical or a linear or branched, optionally at least mono-substituted C₂-C₁₀ alkynyl radical or
 - R⁹ and R¹⁰, together with the bridging nitrogen atom form a saturated or unsaturated, optionally at least mono-substituted, optionally at least one further heteroatom as a ring member containing 5- or 6-membered heterocyclic ring, which may be condensed with a saturated or unsaturated, optionally at least mono-substituted, optionally at least one heteroatom as a ring member containing mono- or bicyclic cycloaliphatic ring system whereby the rings of the ring system are 5- 6- or 7-membered.
- 7.- (Previously Presented) The compound according to claim 6, wherein R⁹ and R¹⁰, identical or different, each represent hydrogen or a linear or branched C₁-C₁₀ alkyl radical, or
 - R⁹ and R¹⁰, together with the bridging nitrogen atom form a radical chosen from a group consisting of

wherein R²⁰, if present, is hydrogen, a linear or branched C₁-C₆ alkyl radical or a benzyl radical.

8.- (Previously Presented) The compound according to claim 1, wherein A and B, identical or different, each represent a linear or branched C₁.C₆ alkyl radical, a linear or branched C₂.C₆ alkenyl radical or a linear or branched C₂.C₆ alkynyl radical, or

A and B, together with the carbon atom to which they are bonded, form a saturated or unsaturated, but not aromatic, optionally at least mono-substituted cycloalkyl ring.

- 9.- (Previously Presented) The compound to claim 1, which is selected from a group consisting of
 - [1] 1-Cyclohexanesulfonyl-3-(1-methyl-1,2,3,6-tetrahydropyridine-4-yl)-5-nitro-1H-indole,
 - [2] 5-Chloro-1-cyclohexanesulfonyl-3-(1-methyl-1,2,3,6-tetrahydropyridine-4-yl)-1H-indole,
 - [3] 5-Amino-1-cyclohexanesulfonyl-3-(1-methyl-1,2,3,6-tetrahydropyridine-4-yl)-1H-indole,
 - [4] 1-Cyclohexanesulfonyl-5-fluoro-3-(1,2,3,5,8,8a-hexahydro-indolizine-7-yl)-1H-indole hydrochloride,
 - a salt thereof, and a solvate thereof.
- 10.- (Previously Presented) A sulfonamide compound of general formula (Ib),

R5
$$R4$$

$$R3$$

$$O = S$$

$$A$$

$$B$$
(Ib)

wherein

R¹ is a -NR⁷R⁸ radical,

R², R³, R⁴, R⁵ and R⁶, identical or different, each represent hydrogen, halogen, cyano, nitro, a saturated or unsaturated, linear or branched aliphatic radical, a linear or branched alkoxy radical, a linear or branched alkylthio radical, hydroxy, trifluoromethyl, a saturated or unsaturated cycloaliphatic radical, an alkylcarbonyl radical, a phenylcarbonyl or a –NR⁹R¹⁰ group,

 R^7 and R^8 , identical or different, each represent hydrogen or a saturated or unsaturated, optionally at least mono-substituted linear or branched C_{1-4} aliphatic radical,

R⁹ and R¹⁰, identical or different, each represent hydrogen or a saturated or unsaturated, linear or branched, optionally at least mono-substituted aliphatic radical, or

R⁹ and R¹⁰, together with the bridging nitrogen atom form a saturated or unsaturated, optionally at least mono-substituted, optionally at least one further heteroatom as a ring member containing heterocyclic ring which may be condensed with a saturated or unsaturated, optionally at least mono-substituted, optionally at least one heteroatom as a ring member containing mono- or bicyclic cycloaliphatic ring system,

A and B, identical or different, each represent a saturated or unsaturated, linear or branched, optionally at least mono-substituted aliphatic radical

or

A and B, together with the carbon atom to which they are bonded, form a saturated or unsaturated, but not aromatic, optionally at least mono-substituted cycloalkyl ring,

and

n is 0;

a stereoisomer thereof, an enantiomer thereof, a diasteromer thereof, a racemate thereof, a pharmaceutically acceptable salt thereof, or mixtures thereof.

- 11.- (Previously Presented) The compound according to claim 10, wherein R², R³, R⁴, R⁵ and R⁶, identical or different, each represent hydrogen, F, Cl, Br, cyano, nitro, a linear or branched C₁.C₆ alkyl radical, a linear or branched C₂.C₆ alkenyl radical, a linear or branched C₁.C₆-alkoxy, a linear or branched C₁.C₆-alkylthio, hydroxy, trifluoromethyl, a saturated or unsaturated C₃.C₈ cycloaliphatic radical, a linear or branched C₁.C₆-alkylcarbonyl radical, phenylcarbonyl or an -NR⁹R¹⁰ group.
- 12.- (Previously Presented) The compound according to claim 10, wherein R⁷ and R⁸, identical or different, wherein R⁷ and R⁸, identical or different, each represent hydrogen, a linear or branched, optionally at least mono-substituted C₁.C₄ alkyl radical with the proviso that R⁷ and R⁸ are not hydrogen at the same time.
- 13.- (Previously Presented) The compound according to claim 10, characterized in that R⁹ and R¹⁰, identical or different, each represent hydrogen, a linear or branched, optionally at least mono-substituted C₁-C₁₀ alkyl radical, a linear or branched, optionally at least mono-substituted C₂-C₁₀ alkenyl radical, or a linear or branched, optionally at least mono-substituted C₂-C₁₀ alkynyl radical or

R⁹ and R¹⁰, together with the bridging nitrogen atom form a saturated or unsaturated, optionally at least mono-substituted, optionally at least one further heteroatom as a ring member containing 5- or 6-membered heterocyclic which may be condensed with a saturated or unsaturated, optionally at least mono-substituted, optionally at least one heteroatom as a ring member containing mono- or bicyclicc cycloaliphatic ring system, whereby the rings of the ring system are 5- 6- or 7-membered.

14.- (Previously Presented) The compound according to claim 13, wherein R⁹ and R¹⁰, identical or different, each represent hydrogen or a linear or branched C₁-C₁₀ alkyl radical, or

R⁹ and R¹⁰, together with the bridging nitrogen atom form a radical chosen from a group consisting of

$$-N$$
 $N-R^{20}$, $-N$ and $-N$

wherein R²⁰, if present, represents hydrogen, a linear or branched C₁-C₆ alkyl radical or a benzyl radical.

- 15.- (Previously Presented) The compound according to claim 10, wherein A and B, together with the carbon atom to which they are bonded, form a saturated or unsaturated, but not aromatic, optionally at least mono-substituted cycloalkyl ring.
- 16.- (Previously Presented) A process for obtaining a sulfonamide compound of general formula (Ia) according to claim 1, wherein at least one compound of general formula (II), or a protected compound thereof,

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wherein A and B have the meaning according to claim 1 and X is a leaving group, is reacted with at least one substituted indole of general formula (III)

wherein R¹-R⁶ and n have the meaning according to claim 1, or a protected compound thereof, and, if necessary, the protective groups are removed.

- 17.- (Previously Presented) A process for obtaining a sulfonamide compound of general formula (Ia) according to claim 1, wherein one or more substituents R², R³, R⁴, R⁵ or R⁶ represent a nitro group, and wherein a sulfonamide compound of general formula (Ia) is reduced to a sulfonamide compound of corresponding general formula (Ia), wherein one or more substituents R², R³, R⁴, R⁵ or R⁶ represent an amino group.
- 18.- (Previously Presented) A process for preparing a salt of the compound of formula (Ia) according to claim 1, the process comprising reacting at least one compound of the general formula (Ia) with a mineral acid or organic acid in a solvent to form the salt of the compound of formula (Ia).
- 19.- (Previously Presented) A composition comprising at least one compound according to claim 1 and one or more pharmacologically acceptable excipients.

20-21-(Cancelled).

- 22.- (Previously Presented) A method of treating a disorder or disease related to food intake in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat the disorder or disease in the subject.
- 23.- (Previously Presented) A method for regulating appetite in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to regulate appetite in the subject.
- 24.- (Previously Presented) A method for regulating body weight in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to regulate body weight in the subject.
- 25.- (Previously Presented) A method of treating obesity in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat obesity in the subject.
- 26.- (Previously Presented) A method of treating bulimia in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat bulimia in the subject.
- 27.- (Previously Presented) A method for treating anorexia in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat anorexia in the subject.

- 28.- (Previously Presented) A method for treating cachexia in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat cachexia in the subject.
- 29.- (Previously Presented) A method for treating type II diabetes in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat type II diabetes in the subject.
- 30.- (Previously Presented) A method of treating a gastrointestinal tract disorder in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat the disorder in the subject.
- 31.- (Previously Presented) A method for treating irritable bowel syndrome in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat irritable bowel syndrome in the subject.
- 32.- (Previously Presented) A method for treating anxiety in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat anxiety in the subject.
- 33.- (Previously Presented) A method for treating depression in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat depression in the subject.

34.- (Previously Presented) A method for treating bipolar disorder in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat the disorder in the subject.

35-42 (Cancelled)

43.- (Previously Presented) A method for treating infantile hyperkinesia in a subject in need thereof, the method comprising administering at least one compound according to claim 1 in an amount sufficient to treat infantile hyperkinesia in the subject.

44-49 (Cancelled)

- 50.- (Previously Presented) A method of treating a disorder or disease related to food intake in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat the disorder or disease in the subject.
- 51.- (Previously Presented) A method for regulating appetite in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to regulate appetite in the subject.
- 52.- (Previously Presented) A method for regulating body weight in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to regulate body weight in the subject.

- 53.- (Previously Presented) A method of treating obesity in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat obesity in the subject.
- 54.- (Previously Presented) A method of treating bulimia in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat bulimia in the subject.
- 55.- (Previously Presented) A method for treating anorexia in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat anorexia in the subject.
- 56.- (Previously Presented) A method for treating cachexia in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat cachexia in the subject.
- 57.- (Previously Presented) A method for treating type II diabetes in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat type II diabetes in the subject.
- 58.- (Previously Presented) A method of treating a gastrointestinal tract disorder in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat the disorder in the subject.

- 59.- (Previously Presented) A method for treating irritable bowel syndrome in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat irritable bowel syndrome in the subject.
- 60.- (Previously Presented) A method for treating anxiety in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat anxiety in the subject.
- 61.- (Previously Presented) A method for treating depression in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat depression in the subject.
- 62.- (Previously Presented) A method for treating bipolar disorder in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat the disorder in the subject.

63-70 (Cancelled).

- 71.- (Previously Presented) A method for treating infantile hyperkinesia in a subject in need thereof, the method comprising administering at least one compound according to claim 10 in an amount sufficient to treat infantile hyperkinesia in the subject.
- 72-74 (Cancelled).
- 75. (Previously Presented) The compound according to claim 1, wherein R¹ represents a NR⁷R⁸ radical or a radical chosen from the group consisting of

wherein, if present, the dotted line represents an optional chemical bond, and R^{19} represents hydrogen, a linear or branched C_1 - C_6 alkyl radical or a benzyl radical, preferably hydrogen or a C_1 - C_2 alkyl radical.

76. (Previously Presented) The compound according to claim 1, wherein R^2 , R^3 , R^4 , R^5 and R^6 , identical or different, each represent H, F, Cl, NO₂, NH₂ or a C₁₋₂ alkyl radical.